

Variations in the GLA gene correlate with globotriaosyl-  
globotriaosylsphingosine analog levels in urine and plas

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Tandem Mass Spectrometry Quantitation of Lyso-Gb 3 and Six Related Analogs in Plasma for Fabry Disease Patients. <i>Current Protocols in Human Genetics</i> , 2016, 90, 17.23.1-17.23.9.	3.5	14
2	Relative distribution of Gb <sub>3</sub> isoforms/analogs in NOD/SCID/Fabry mice tissues determined by tandem mass spectrometry. <i>Bioanalysis</i> , 2016, 8, 1793-1807.	0.6	14
3	Tandem Mass Spectrometry of Sphingolipids. <i>Advances in Clinical Chemistry</i> , 2016, 77, 177-219.	1.8	5
4	Biomarkers associated with clinical manifestations in Fabry disease patients with a late-onset cardiac variant mutation. <i>Clinica Chimica Acta</i> , 2017, 466, 185-193.	0.5	44
5	Characterization of Classical and Nonclassical Fabry Disease: A Multicenter Study. <i>Journal of the American Society of Nephrology: JASN</i> , 2017, 28, 1631-1641.	3.0	244
6	A simple method for quantification of plasma globotriaosylsphingosine: Utility for Fabry disease. <i>Molecular Genetics and Metabolism</i> , 2017, 122, 121-125.	0.5	24
8	European expert consensus statement on therapeutic goals in Fabry disease. <i>Molecular Genetics and Metabolism</i> , 2018, 124, 189-203.	0.5	122
9	&p&gt;Multiple phenotypic domains of Fabry disease and their relevance for establishing genotype&ndash;phenotype correlations&lt;p&gt;. <i>The Application of Clinical Genetics</i> , 2019, Volume 12, 35-50.	1.4	37
10	Effectiveness of plasma lyso-Gb3 as a biomarker for selecting high-risk patients with Fabry disease from multispecialty clinics for genetic analysis. <i>Genetics in Medicine</i> , 2019, 21, 44-52.	1.1	61
11	Biomarkers in Andersonâ€“Fabry Disease. <i>International Journal of Molecular Sciences</i> , 2020, 21, 8080.	1.8	35
12	A comprehensive testing algorithm for the diagnosis of Fabry disease in males and females. <i>Molecular Genetics and Metabolism</i> , 2020, 130, 209-214.	0.5	26
13	Novel biomarkers for lysosomal storage disorders: Metabolomic and proteomic approaches. <i>Clinica Chimica Acta</i> , 2020, 509, 195-209.	0.5	5
14	An expert consensus document on the management of cardiovascular manifestations of Fabry disease. <i>European Journal of Heart Failure</i> , 2020, 22, 1076-1096.	2.9	96
15	Circular RNA-based biomarkers in blood of patients with Fabry disease and related phenotypes. <i>Journal of Medical Genetics</i> , 2021, , jmedgenet-2020-107086.	1.5	2
16	Biomarkers in Fabry Disease. Implications for Clinical Diagnosis and Follow-up. <i>Journal of Clinical Medicine</i> , 2021, 10, 1664.	1.0	12
17	Narrative review on Morbus Fabry: diagnosis and management of cardiac manifestations. <i>Cardiovascular Diagnosis and Therapy</i> , 2021, 11, 650-660.	0.7	5
18	Plasma Globotriaosylsphingosine and Î±-Galactosidase A Activity as a Combined Screening Biomarker for Fabry Disease in a Large Japanese Cohort. <i>Current Issues in Molecular Biology</i> , 2021, 43, 389-404.	1.0	2
19	Clinical and diagnostic aspects of Fabry disease management: a narrative review with a particular focus on Brazilian expertsâ€™ perspectives. <i>Journal of Inborn Errors of Metabolism and Screening</i> , 0, 10, .	0.3	1

#	ARTICLE	IF	CITATIONS
20	Metabolic Fingerprinting of Fabry Disease: Diagnostic and Prognostic Aspects. <i>Metabolites</i> , 2022, 12, 703.	1.3	4
21	Late-onset and classic phenotypes of Fabry disease in males with the <i>GLA</i> -Thr410Ala mutation. <i>Open Heart</i> , 2023, 10, e002251.	0.9	1